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(74) Agents: WOLFF, Jessica, R. et al.; Lyon & Lyon LLP, Suite 4700, 633 West Fifth Street, Los Angeles, CA 90071-2066

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(71) Applicants (for all designated States except US): GLYCOMED INCORPORATED [US/US]; c/o Ligand Pharmaceuticals Incorporated, 10275 Science Center Drive, San Diego, CA 92121 (US). SANKYO CO., LTD. [JP/JP]; 2-58, Hiromachi 1-chome, Shinagawa-ku, Tokyo 140-8710 (JP).

(72) Inventors; and

(75) Inventors/Applicants (for US only): ANDERSON, Mark, B. [US/US]; 41 Las Cascadas Road, Orinda, CA 94563 (US). LEVY, Daniel, E. [US/US]; 3918 Christian Drive, Belmont, CA 94002 (US). HOLME, Kevin, R. [US/US]; 13644 Landfair Road, San Diego, CA 92130 (US).

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(57) Abstract

The present invention discloses medicaments that are selectin-ligand structural mimetics that bind to certain selectins wherein the mimetics may lack the sialic acid and/or fucose of the natural selecting ligand, sialyl Lewis* (sLe*), but have a structure capable of mimicking the structural features necessary for selectin recognition. In particular, the invention compounds mimic the key structural features of the oligosaccharides responsible for selectin-mediated cell adhesion. These features consist of the charge-distance-coordination relationship between the carboxylic acid functionality of sialic acid at a distance of 8-12 angstroms of the L-fucose moiety. The invention compounds are disalicylate, its analogs, and disalicylate-based C-glycoside compounds. The present invention also discloses methods of treating selectin-mediated disorders comprising administering the compounds disclosed.

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